

# ● PRINTER RUSH ●

(PTO ASSISTANCE)

IFW

Application: <u>09535364</u>	Examiner: <u>CANELLA</u>	GAU: <u>1642</u>
From: <u>IF</u>	Location: <u>UDC</u> FMF FDC	Date: <u>8-23-05</u>

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DOC CODE	DOC DATE	MISCELLANEOUS
<input type="checkbox"/> 1449	_____	<input type="checkbox"/> Continuing Data
<input type="checkbox"/> IDS	_____	<input checked="" type="checkbox"/> Foreign Priority
<input type="checkbox"/> CLM	_____	<input type="checkbox"/> Document Legibility
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[RUSH] MESSAGE: A 35 U.S.C. 119(a)-(d)  
foreign priority claim cannot  
be based on a U.S. application.  
Please make all necessary corrections  
to file wrapper & specifications.  
See MPEP 1293.03(c) priority under  
35 U.S.C. 120.  
Thank you

[XRUSH] RESPONSE: A PCT application can be used as a  
foreign priority document. Please see the attached examples  
of recently issued patents: US 6,884,771, US 6,887,974,  
US 6,855,559, US 6,800,604, US 6,790,624.

INITIALS: KAL

NOTE: This form will be included as part of the official USPTO record, with the Response document coded as XRUSH.  
 REV 10/04



US006884771B1

(12) **United States Patent**  
**Acton et al.**

(10) **Patent No.:** **US 6,884,771 B1**  
**(45) Date of Patent:** **Apr. 26, 2005**

(54) **ANGIOTENSIN CONVERTING ENZYME  
HOMOLOG AND USES THEREFOR**

(75) **Inventors:** Susan Acton, Lexington, MA (US);  
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(US)

(73) **Assignee:** Millennium Pharmaceuticals, Inc.,  
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(\*) **Notice:** Subject to any disclaimer, the term of this  
patent is extended or adjusted under 35  
U.S.C. 154(b) by 472 days.

(21) **Appl. No.:** 09/635,501

(22) **Filed:** Aug. 9, 2000

**Related U.S. Application Data**

(63) Continuation-in-part of application No. 09/407,427, filed on  
Sep. 29, 1999, which is a continuation-in-part of application  
No. 09/163,648, filed on Sep. 30, 1998, which is a continu-  
ation-in-part of application No. 08/989,299, filed on Dec. 11,  
1997.

(30) **Foreign Application Priority Data**

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(51) **Int. Cl.<sup>7</sup>** ..... A51K 38/00

(52) **U.S. Cl.** ..... 514/2; 514/12; 530/350;  
530/361; 424/94.1; 424/94.6; 435/183;  
435/195

(58) **Field of Search** ..... 514/2, 12; 530/350,  
530/361; 424/94.1, 94.6; 435/183, 195;  
536/23.5; 800/7

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*Assistant Examiner*—B. Dell Chism

(74) *Attorney, Agent, or Firm*—Millennium  
Pharmaceuticals, Inc.

(57) **ABSTRACT**

The present invention relates to the discovery of novel genes  
encoding an angiotensin converting enzyme, Angiotensin  
Converting Enzyme-2 (ACE-2). The invention provides  
therapeutics, prognostic and diagnostics methods for treat-  
ing blood pressure related disorders as well as various types  
of allergic conditions, among others. Also disclosed are  
screening assays for identifying compounds for treating and  
preventing these conditions.

**25 Claims, 23 Drawing Sheets**



US006887974B2

(12) **United States Patent**  
**Pathak**(10) **Patent No.:** **US 6,887,974 B2**  
(45) **Date of Patent:** **May 3, 2005**(54) **CROSSLINKING AGENTS AND METHODS OF USE**(75) **Inventor:** Chandrashekhar P. Pathak, Austin, TX (US)(73) **Assignee:** Incept LLC, Lexington, MA (US)(\*) **Notice:** Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 421 days.(21) **Appl. No.:** 10/068,807(22) **Filed:** Feb. 5, 2002(65) **Prior Publication Data**

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**Related U.S. Application Data**

(62) Division of application No. 09/147,897, filed on Aug. 30, 1999.

(60) Provisional application No. 60/026,526, filed on Sep. 23, 1996, provisional application No. 60/039,904, filed on Mar. 4, 1997, and provisional application No. 60/040,417, filed on Mar. 13, 1997.

(30) **Foreign Application Priority Data**

Sep. 22, 1997 (WO) ..... PCT/US97/16897

(51) **Int. Cl.<sup>7</sup>** ..... C09F 15/00; C08G 63/48; C08G 63/08(52) **U.S. Cl.** ..... 530/200; 530/350; 530/380; 530/382; 525/54.1; 525/54.11; 525/54.2; 528/354; 528/361(58) **Field of Search** ..... 530/200, 350, 530/380, 382; 525/54.1, 54.11, 54.2; 528/354, 361(56) **References Cited****U.S. PATENT DOCUMENTS**

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**Primary Examiner**—Jon Weber**Assistant Examiner**—Abdel A. Mohamed(74) **Attorney, Agent, or Firm**—Patterson, Thunte, Skaar & Christensen, P.A.(57) **ABSTRACT**

Polymeric crosslinking agents are disclosed that have an inert water soluble polymeric component, biodegradable components, functional components reactive with chemical groups on a protein, for example, amine or thiol groups. The inert polymeric component may be flanked at each end with a biodegradable component which is flanked at each end with a protein reactive functional component. A polymeric crosslinking agent is disclosed having a biodegradable component, polyalkylene oxide, and at least three reactive functional groups that are each capable of forming a covalent bond in water with at least one functional group such as an amine, thiol, or carboxylic acid.

**25 Claims, 7 Drawing Sheets**



US00685559B1

(12) **United States Patent**  
Christensen et al.

(10) Patent No.: **US 6,855,559 B1**  
(45) Date of Patent: **Feb. 15, 2005**

(54) **REMOVAL OF EMBEDDING MEDIA FROM BIOLOGICAL SAMPLES AND CELL CONDITIONING ON AUTOMATED STAINING INSTRUMENTS**

(75) Inventors: **Kimberly Christensen**, Tucson, AZ (US); **Ethel R. Macrea**, Tucson, AZ (US); **Noemi Sebastiao**, Tucson, AZ (US)

(73) Assignee: **Ventana Medical Systems, Inc.**, Tucson, AZ (US)

(\*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 81 days.

(21) Appl. No.: **09/721,096**

(22) Filed: **Nov. 22, 2000**

#### Related U.S. Application Data

(60) Provisional application No. 60/099,018, filed on Sep. 3, 1998.

#### Foreign Application Priority Data

Feb. 26, 1999 (WO) ..... PCT/US99/04181  
Sep. 3, 1999 (WO) ..... PCT/US99/20353

(51) Int. Cl.<sup>7</sup> ..... **G01N 1/18**

(52) U.S. Cl. .... **436/177; 436/174; 436/175; 436/139**

(58) Field of Search ..... **436/139, 174-177; 422/61, 100, 58; 427/2.11; 455/40.52, 40.5**

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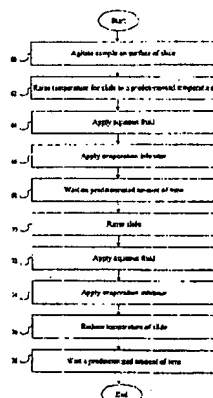
Primary Examiner—Yelena G. Gakh

(74) Attorney, Agent, or Firm—McDonnell Boehnen Hulbert & Berghoff LLP

#### (57) ABSTRACT

The present invention provides reagents for use in an automated environment for removing or etching embedding media by exposing a biological sample to be stained in histochemical or cytochemical procedures without the dependence on organic solvents. The reagents comprise components optimized to facilitate removal or etching of the embedding media from the biological sample. The present invention also provides reagents for use in an automated environment for cell conditioning biological samples wherein the cells are predisposed for access by reagent molecules for histochemical and cytochemical staining procedures. The reagents comprise components optimized to facilitate molecular access to cells and cell constituents within the biological sample.

16 Claims, 8 Drawing Sheets





US006800604B2

(12) **United States Patent**  
Gurney et al.

(10) Patent No.: **US 6,800,604 B2**  
(45) Date of Patent: **Oct. 5, 2004**

(54) **POLYPEPTIDES THAT INHIBIT HUMAN  
SERUM-INDUCED CLEAVAGE OF  
HEPATOCTE GROWTH FACTOR**

(75) Inventors: **Austin L. Gurney**, Belmont, CA (US);  
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(73) Assignee: **Genentech, Inc.**, South San Francisco,  
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(\*) Notice: Subject to any disclaimer, the term of this  
patent is extended or adjusted under 35  
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(21) Appl. No.: **09/742,201**

(22) Filed: **Dec. 19, 2000**

(65) **Prior Publication Data**

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**Related U.S. Application Data**

(60) Provisional application No. 60/253,665, filed on Nov. 28,  
2000.

(30) **Foreign Application Priority Data**

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(51) Int. Cl.<sup>7</sup> ..... **C07K 14/00; A61K 38/00**

(52) U.S. Cl. .... **514/2; 530/300; 530/350;**  
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(58) Field of Search ..... **530/300, 350,**  
**530/351; 514/2, 12; 424/85.1, 184.1, 192.1,**  
**198.1**

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Primary Examiner—Elizabeth Kemmerer

Assistant Examiner—Bridget E. Bunner

(74) Attorney, Agent, or Firm—Paul Naik; Craig Svoboda

(57)

**ABSTRACT**

Compositions and methods are disclosed for stimulating or  
inhibiting angiogenesis and/or cardiovascularization in  
mammals, including humans. Pharmaceutical compositions  
are based on polypeptides or antagonists thereto that have  
been identified for one or more of these uses. Disorders that  
can be diagnosed, prevented, or treated by the compositions  
herein include trauma such as wounds, various cancers, and  
disorders of the vessels including atherosclerosis and cardiac  
hypertrophy. In addition, the present invention is directed to  
novel polypeptides and to nucleic acid molecules encoding  
those polypeptides. Also provided herein are vectors and  
host cell comprising those nucleic acid sequences, chimeric  
polypeptide molecules comprising the polypeptides of the  
present invention fused to heterologous polypeptide  
sequences, antibodies which bind to the polypeptides of the  
present invention and to methods for producing the polypep-  
tides of the present invention.

**27 Claims, 5 Drawing Sheets**



US006790624B2

(12) **United States Patent**  
Mayer

(10) **Patent No.:** US 6,790,624 B2  
(45) **Date of Patent:** Sep. 14, 2004

(54) **COILED-COIL MEDIATED  
HETERODIMERIZATION FUNCTIONAL  
INTERACTION TRAP**

(75) **Inventor:** Bruce J. Mayer, Tolland, CT (US)

(73) **Assignee:** The University of Connecticut,  
Farmington, CT (US)

(\*) **Notice:** Subject to any disclaimer, the term of this  
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(21) **Appl. No.:** 09/816,756

(22) **Filed:** Mar. 24, 2001

(65) **Prior Publication Data**

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**Related U.S. Application Data**

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1999.

(30) **Foreign Application Priority Data**

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(51) **Int. Cl.<sup>7</sup>** ..... G01N 33/53

(52) **U.S. Cl.** ..... 435/7.1; 435/6; 435/5;  
435/4; 530/350; 536/23.4

(58) **Field of Search** ..... 435/7.1, 6, 5, 4;  
530/350; 324; 536/23.4

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(57) **ABSTRACT**

Fusion proteins containing coiled-coil heterodimerization  
domains substituted for modular protein binding domains  
useful for validating functionally relevant protein-protein  
interactions, directing enzymes to specific substrates, and  
screening fusion libraries for functionally important inter-  
action partners.

**2 Claims, 3 Drawing Sheets**